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Uppsala, Sweden, *March 1, 2018*  
Ex officio:

*[Signature]*  
Olof Wijk  
Notary Public



The Ministry for Foreign Affairs in Stockholm, hereby certifies that  
**Mr. Olof Wijk**  
Notary Public of Uppsala  
has issued and signed the foregoing attestation in his official capacity.




Stockholm 5 March 2018

Nr. 4856

Fee Paid 180 SEK

*[Signature]*  
Essie Persson

  
CHỨNG NHẬN/HỢP PHÁP HÓA LÃNH SỰ  
CONSULAR AUTHENTICATION

1. Quốc gia... *Viet Nam / Vietnam*  
Country

2. do Ông (Bà)... *Essie Persson*... ký  
has been signed by

3. với chức danh  
acting in the capacity of

4. và con dấu của *Bộ Ngoại Giao Thụy Điển*  
bears the seal/stamp of *Swedish Ministry for Foreign Affairs*

được chứng nhận/hợp pháp hóa lãnh sự  
Certified

5. tại *Stockholm*  
at

6. ngày *05.02.2018*

7. Cơ quan cấp *ĐSQ VN / VN Embassy*  
by

8. Số: *111/2018*  
Nº

TL. ĐẠI SỨ / FOR THE AMBASSADOR  
Bí thư thứ Hai / Second Secretary

*[Signature]*  
**Phạm Thị Quyên**

**EG-KONFORMITÄTSERKLÄRUNG**  
**EC DECLARATION OF CONFORMITY**

gemäß Anhang III der Richtlinie 98/79/EC über In-vitro-Diagnostika  
 according to Annex III of the IVD Medical Device Directive 98/79/EC

Hersteller / Manufacturer:	Kibion GmbH
Branche / Business sector:	Atemtests / Breath tests
Anschrift / Address:	Haferwende 31 D-28357 Bremen, Germany
IVD-Medizinprodukt / IVD-Medical Device:	Kibion Dynamic Pro
Klassifizierung gemäß 98/79/EG / Classification according to 98/79/EC:	Sonstiges Produkt (nicht im Anhang II, nicht zur Eigenanwendung) Other product (not in Annex II, not for self- testing)
Nomenklatursystem & -Code/ Code and Term:	EDMS 26-09. Kategorie 06

Angaben zum Produkt / Device identification		
Produkt / Product	Artikel Nr./ REF no.	Beschreibung, Version etc. Description, version, etc.
Kibion® Dynamic Pro	8032	Erweiterungseinheit für das Kibion® Dynamic Base (Infrarot Isotopen Analysator für 13C- Atemgas) mit 16 Anschlüssen Port extension unit for IRIS Dynamic base (Infrared Isotope Analyser for 13C breath tests) with 16 ports

Wir erklären in alleiniger Verantwortung, dass die  
Medizinprodukte allen anwendbaren  
Anforderungen der Richtlinie 98/79/EC für in-  
vitro-Diagnostika bzw. des Gesetzes über  
Medizinprodukte (MPG) entsprechen.

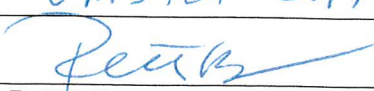
We declare, in sole responsibility, that the  
concerned medical devices comply with the  
applicable requirements of the IVD-directive  
98/79/EC and with the German law concerning  
Medical Devices (MPG) respectively.

Das Produkt ist CE-zertifiziert durch Kibion  
GmbH.

The product is CE-marked by Kibion GmbH.

Nicht von uns ermächtigte Produkt-änderungen  
machen diese Erklärung ungültig.

Any modification of the devices, not authorised  
by us, will invalidate this declaration.

Ort und Datum / Place and Date	Uppsala 2017-03-10
Unterschrift / Signature	
Name / Printed name	Petter Bäckgren
Funktion / Position	Geschäftsführer / CEO

